PCT

WORLD INTELLECTUAL PROPERTY ORGANIZATION International Bureau



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 6:

C10M 105/42, 105/44, 105/46, 129/78, 129/80, 129/82, 169/04, 171/00, C10L 1/18, C07C 69/34, 69/50, 69/593, 69/604

A1

(11) International Publication Number:

WO 99/16849

(43) International Publication Date:

8 April 1999 (08.04.99)

(21) International Application Number:

PCT/EP98/06145

(22) International Filing Date:

28 September 1998 (28.09.98)

(30) Priority Data:

97202992.0

i October 1997 (01.10.97)

EP

(71) Applicant (for all designated States except US): UNICHEMA CHEMIE B.V. [NL/NL]; Buurtje I, NL-2802 BE Gouda (NL).

(72) Inventors; and

- (75) Inventors/Applicants (for US only): KENBEEK, Dirk [NL/NL]; Strengen 5, NL-3421 JS Oudewater (NL). VERBOOM, Comelis [NL/NL]; Ravelplein 47, NL-2807 HJ Gouda (NL). VAN DER WAAL, Gijsbert [NL/NL]; Albert Schweitzerstraat 22, NL-2861 XX Bergambacht (NL).
- (74) Agents: GEARY, Stephen et al.: ICI Group Intellectual Property, P.O. Box 11, The Heath, Runcom, Cheshire WA7 4QE (GB).

(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), Isuropean patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

Published

With international search report.

Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.

(54) Title: COMPLEX ESTERS, FORMULATIONS COMPRISING THESE ESTERS AND USE THEREOF

(57) Abstract

An ester resulting from an esterification reaction between at least one polyfunctional alcohol and at least one polyfunctional carboxylic acid using a chain stopping agent to form ester bonds with the remaining hydroxyl or carboxyl groups is disclosed. The polyfunctional carboxylic acid comprises an aliphatic dicarboxylic acid containing from 9 to 18 carbon atoms, dimerised and/or trimerised fatty acids or mixtures thereof, with the proviso that dimerised and trimerised fatty acids do not constitute more than 80% by weight of the total amount of polyfunctional carboxylic acid used. The chain stopping agent may be a monocarboxylic acid or a monofunctional alcohol having at least 14 carbon atoms. The complex esters have a kinematic viscosity at 100 C of from 30 to 1000 cSt, preferably from 30 to 200 cSt. The complex ester is useful "as is" or as an additive and/or as a base fluid and/or a thickener in transmission oils, hydraulic fluids, four–stroke oils, fuel additives, compressor oils, greases, chain oils and for metal working metal rolling applications. A multigrade gear oil formulation comprising one or more of the above complex esters is also part of the invention.

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AI.	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
\mathbf{AM}	Armenia	Fl	Finland	LT	Lithuania	SK	Słovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
ΑÜ	Australia	GA	Gaben	LV	Latvia	SZ	Swaziland
AZ.	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	gascar TJ	
BE	Belgium	GN	Guinea	MK	The former Yugoslav	TM	Turkmenistan
BF	Burkina Faso	GR	Greece		Republic of Macedonia	TR	Turkey
BG	Bulgaria	пc	Hungary	ML	Mali	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MN	MN Mongolia UA		Ukraine
BR	Brazi!	IL.	Israei	MR	Mauritania	บัต	Uganda
BY	Belanis	18	Iceland	MW	Malawi US		United States of America
CA	Canada	IT.	Italy	MX	Mexico UZ		Uzbekistan
CF	Central African Republic	JР	Japan	NE	Niger	VN	Viet Nam
CG	Congo	KE	Kenya	NL	Netherlands	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NO	Norway	ZW	Zinibahwe
CI	Côte d'Ivoire	КP	Democratic People's	NZ	New Zealand		
CM	Cameroon		Republic of Korea	PL	Poland		
CN	China	KR	Republic of Korea	PT	Portugul		
CU	Cuba	KZ	Kazakstan	RO	Romania		
CZ.	Czech Republic	LC	Saint Lucia	Rt'	Russian Federation		
DE	Germany	1.1	Liechtenstein	SD	Sudan		
DK	Denmark	LK	Sri Lanka	SF.	Sweden		
E.F.	Estonia	LR	Liberia	SG	Singapore		

COMPLEX ESTERS, FORMULATIONS COMPRISING THESE ESTERS AND USE THEREOF

5

10

15

20

25

30

The present invention relates to esters containing more than one ester linkage, hereinafter known as "complex" esters, to formulations comprising one or more of these complex esters and to various uses of the complex esters and the formulations. More specifically, the present invention relates to complex esters and their use as an additive and/or a base fluid and/or thickener in various types of formulations suitable for use in lubrication applications, for example gear oils, hydraulic fluids, compressor oils, greases and four-stroke oils. The present invention also relates to formulations comprising one or more of these complex esters.

Complex esters are known in the art. For instance, DE-A-2620645 discloses a process for lubricating a two stroke engine by using a two stroke lubricating oil of which the base oil consists of at least one hydrocarbon oil and a complex ester. The complex ester results from esterification of trimethylolpropane with at least one saturated. linear or slightly branched C_2 - C_{36} saturated, aliphatic dicarboxylic acid and a mixture of at least one linear or slightly branched C_2 - C_{14} monocarboxylic acid and at least one saturated, linear or slightly branched aliphatic C_{15} - C_{30} monocarboxylic acid. Maximum kinematic viscosity at 98.9 C (Vk, $_{98.9}$) of the complex ester suitably is only 25 cSt, which corresponds to a typical viscosity of a two-stroke oil.

In FR-A-2,187.894, a process for lubricating two stroke engines or rotary engines is disclosed, wherein use is made of a lubricating oil of which the base oil is a complex ester having a kinematic viscosity of more than 6 cSt at 98,9 C. In this reference complex esters are defined as esters formed by condensation of a polycarboxylic acid with a mono- and polyalcohol or as esters formed by condensation of a polyol with a poly- and monocarboxylic acid. Several examples of complex esters are given: adipate/trimethylolpropane/heptanol having a Vk.,98,9 of 19.2 cSt. adipate/trimethylolpropane/dodecanoic acid having a Vk.,98,9 of 13,7 cSt and azelaic acid/pentaerythritol/heptanoic acid/dodecanoic acid having a Vk.,98,9 of 15,4 cSt. Again, these low viscosities are typical for two-stroke engine oils.

DE-A-2130850 discloses a lubricant composition containing or consisting of at least one low viscosity and one high viscosity component, where the high viscosity component is a complex ester having a kinematic viscosity at 99 C of more than 50 cSt.

and a flat viscosity-temperature behaviour. The complex esters are obtained by esterification of unbranched dicarboxylic acids having at least 10 carbon atoms with tri- or tetrafunctional alcohols and stopping with monoalcohols of which at least 25% is linear and low molecular. Trimethylolpropane and pentaerythritol are listed as suitable alcohols, whilst n-butanol and n-hexanol are mentioned as suitable low molecular monoalcohol chain stopping agent.

It has been found that complex esters having improved properties can be obtained by selecting certain compounds for use in the production of the complex ester so as to reduce or remove the number of free alcohol and/or carboxylic acid groups in the ester and so terminate the esterification process. Such compounds are hereinafter referred to as "chain stopping agents". We have found that monoalcohols having relatively long carbon chains, i.e. of 14 carbon atoms or more, or monocarboxylic acids having at least 7 carbon atoms provide surprising improvements in properties of the complex esters.

10

15

20

25

30

In WO-A-97/08277 two categories of ester base stocks for smokeless two stroke engine lubricants are disclosed. The first category are ester base stocks comprising a first ester having a viscosity at 100 C of 2 cSt or less and a second ester having a viscosity such that when the first and second ester are mixed, the resulting mixture has a viscosity at 100 C of 3.0 to 20.0 cSt and a smoke index of at least 75. The second ester may be a stopped, i.e. chain terminated, or unstopped, i.e. still having some functionality, complex ester. The second category of ester base stocks is formed by one or more esters selected from the group consisting of (a) linear oligoesters having a molecular weight of 3000 Daltons or less, (b) complex, non-hindered polyesters wherein the polyol is a molecule having one or more beta hydrogen atoms. (c) complex. non-hindered polyesters wherein the polyol component is a non-hindered polyol having at least 3 OH groups and (d) esters wherein the polyol component is a hindered polyol and the carboxylic acid is a mono- or polycarboxylic acid or a mixture thereof. Several complex esters of the various categories are described, but most of them have a relatively low kinematic viscosity. The stopped complex ester having the highest kinematic viscosity at 100 C (44.5 cSt) is an ester of trimethylolpropane, dimer acid and oleic acid (C18:1 monoacid) as the chain stopping agent.

However, it has been found that the use of dimer acid, i.e. mainly dimerised fatty acids also comprising some trimerised fatty acids, as the sole polycarboxylic acid component has some disadvantages in terms of interaction with certain additive packages comprising sulphur- and/or phosphorus-containing components. Therefore, it would be advantageous to provide a complex ester not comprising dimer acid as the sole polycarboxylic acid component. Furthermore, it would be advantageous if such stopped complex esters could be provided having high kinematic viscosities at 100 C, i.e. 30 cSt or higher.

5

10

15

25

30

The present invention aims to provide a complex ester having a relatively high viscosity, which can be used as a functional fluid itself or in various formulations as a functional fluid, for example a lubricating formulation. Furthermore, and depending on the application, the complex ester should provide high oxidation stability and excellent lubricity, whilst, desirably, possessing good biodegradability characteristics. It will be appreciated that the latter is highly desired in view of the increasing environmental awareness and corresponding demand for environmentally friendly products.

Accordingly, the first aspect of the invention relates to a complex ester obtainable by an esterification reaction between at least one polyfunctional alcohol and at least one polyfunctional carboxylic acid and a chain stopping agent, wherein

- 20 (a) the polyfunctional alcohol is a hindered or non-hindered, aliphatic polyol,
 - (b) the polyfunctional carboxylic acid comprises an aliphatic dicarboxylic acid containing from 9 to 18 carbon atoms, dimerised and/or trimerised fatty acids or mixtures thereof, with the proviso that dimerised and trimerised fatty acids do not constitute more than 80% by weight, preferably not more than 50% by weight, of the total amount of polyfunctional carboxylic acid used.
 - (c) the chain stopping agent comprises either an aliphatic monocarboxylic acid selected from the group consisting of straight chain saturated acids containing from 7 to 22, preferably from 7 to 14, carbon atoms, branched saturated acids containing from 7 to 24 carbon atoms, straight or branched unsaturated acids containing from 16 to 24 carbon atoms and mixtures thereof or at least one aliphatic, straight or branched, saturated or unsaturated.

monofunctional alcohol containing at least 14 carbon atoms, and preferably not having more than 24 carbon atoms, and

(d) the complex ester has a kinematic viscosity at 100 C (Vk,₁₀₀) of from 30 to 1000 cSt, preferably from 30 to 200 cSt.

5

10

15

20

25

30

Preferably the complex ester according to the first aspect of the invention is obtained by an esterification reaction between at least one polyfunctional alcohol and at least one polyfunctional carboxylic acid and a chain stopping agent

The polyfunctional alcohol preferably is a hindered polyol, more preferably a neopentyl polyol. Examples of suitable neopentyl polyols are neopentyl glycol, dipentaerythritol, trimethylolpropane and pentaerythritol, the latter two being particularly preferred.

The polyfunctional carboxylic acid preferably comprises at least one aliphatic dicarboxylic acid having from 9 to 12 carbon atoms, more preferably selected from nonanedioic acid, decanedioic acid, dodecanedioic acid and mixtures thereof. The presence of dimerised and/or trimerised fatty acids is also considered beneficial provided the amount of such acids does not exceed 80% by weight, preferably 50% by weight, of the total amount of polyfunctional carboxylic acids used. Dimerised and/or trimerised fatty acids may be obtained by subjecting an unsaturated fatty acid-containing feedstock to dimerisation by heat treatment in the presence of a suitable catalyst, as is well known in the art. Suitable unsaturated fatty acid containing sources usually comprise a mixture of unsaturated fatty acids with oleic acid (C18:1) often being the main component beside other mono- and polyunsaturated fatty acids. Dimer acid ("C36di") is produced in substantial quantities in the dimerisation reaction. The final product, which is used for manufacturing the complex esters of the invention. usually is a mixture of dimers and trimers commonly in a dimer/trimer ratio of about 80/20. This mixture contains aliphatic as well as cyclic structures including both naphthenic and aromatic structures. If desired, dimers and/or trimers of high purity (e.g. 95% or more) can be manufactured by molecular distillation of the aforementioned mixture of dimers and trimers. This mixture of dimers and trimers as well as purified dimers and/or trimers can be used as the dimerised and/or trimerised fatty acid component. If desired, the dimerised and/or trimerised fatty acid(s) used can be subjected to hydrogenation prior to being used for forming the complex ester.

Suitably, the polyfunctional carboxylic acid is not dimerised and/or trimerised acid alone, as it was found that this may affect the oxidation performance of for instance a gear oil formulation. It was found that a maximum level of 80% by weight of dimerised and/or trimerised acid, based on total weight of polyfunctional carboxylic acid used, still results in an acceptable oxidation stability. The best results are, however, attained when the dimerised and/or trimerised acid does not constitute more than 50% by weight, more preferably not more than 35% by weight, of the total amount of polyfunctional carboxylic acid used.

5

10

15

20

25

30

The chain stopping agent is used to react with the reactive OH- or COOH-groups, as may be the case, which remain unreacted after reaction between the polyfunctional alcohol and the polyfunctional carboxylic acid. The chain stopping agent should preferably have a relatively long carbon chain for achieving optimum viscosity properties (i.e. a kinematic viscosity at 100 C of at least 30 cSt). In those applications where oxidation stability is very important, such as in gear oil formulations, the chain stopping agent preferably should be saturated. For applications where oxidation stability is less critical, such as for instance in hydraulic fluids, unsaturated fatty acids like olein (technical grade eleic acid) or unsaturated alcohols may also be used. Of the chain stopping agents mentioned above, isostearic acid (isoC18) is very much preferred. However, other fatty acids, like palmitic acid (C16) or stearic acid (C18) are also useful. Furthermore, monocarboxylic acids such as octanoic acid and decanoic acid can also be used. Guerbet acids are also included among the suitable monocarboxylic acids. Examples of suitable monofunctional alcohols are tetradecanol, isotetradecanol, octadecanol and iso-octadecanol. Guerbet alcohols are also included among the suitable monofunctional alcohols.

The complex ester according to the present invention should have a Vk_{100} of from 30 to 1000 cSt and preferably from 30 to 200 cSt. For certain applications, such as in gear oils, it is preferred that the Vk_{100} has a value of from 100 to 140 cSt. The kinematic viscosity at 40 C (Vk_{140}) of the complex esters suitably has a value in the range of from 230 to 20,000 cSt, more suitably from 230 to 2800 cSt.

The polyol, polyfunctional carboxylic acid(s) and chain stopping agent, which react to form the complex ester, are preferably used in the following amounts depending in the specific materials employed ("pbw" are parts by weight):

15-20 pbw of polyol,

5

10

15

20

25

30

20-25 pbw polyfunctional carboxylic acid and

55-65 pbw chain stopping agent.

The materials are selected so as to provide a complex ester having a Vk,₁₀₀ within the preferred range of from 100 to 140 cSt.

The complex ester according to the present invention can suitably be used in combination with an extreme pressure and/or anti-wear additive (hereinafter EP/AW) containing sulphur and/or phosphorus-containing compounds e.g. in gear oils.

Accordingly, a further aspect of the invention invention relates to a formulation comprising a complex ester as described according to the first aspect of the invention and a sulphur and/or phosphorus-containing EP/AW additive package in a weight ratio of complex ester to additive package of from 1:3 to 9:1. Suitable sulphur and/or phosphorus-containing EP/AW additive packages are well known in the art, particularly for use in gear oils to avoid wear of the gear wheels. Commercially available sulphur-phosphorus-containing EP/AW additive packages are, for instance, manufactured by Ethyl Corporation, Lubrizol and Paramins.

The complex ester according to the invention can be used as a functional fluid in many different applications, for example in lubricating formulations. The ester may be used as a functional fluid or as an additive and/or a base fluid and/or as a thickener in a functional fluid composition.

Thus, the present invention also relates to the use of the complex ester described according to the first aspect of the invention as a functional fluid.

The present invention also relates to functional fluid compositions comprising the complex ester described according to the first aspect of the invention.

The invention also relates to the use of a formulation containing the complex ester as described in the first aspect of the invention as functional fluid composition, such as transmission oils, for example automotive and industrial gear oils, axle oils and automatic transmission fluids, and also in hydraulic fluids, four-stroke oils, fuel additives, compressor oils, greases, chain oils and for metal working and metal rolling applications.

Examples of functional fluids and functional fluid compositions include transmission oils, for example automotive and industrial gear oils, axle oils and

automatic transmission fluids, and also in hydraulic fluids, four-stroke oils, fuel additives, compressor oils, greases, chain oils and for metal working and metal rolling applications.

It has been found that the complex ester according to the invention is particularly suitable to be used as a high viscosity base fluid and/or a thickener in multigrade gear oil formulations.

5

10

15

20

25

30

Multigrade gear oil formulations comprising a synthetic thickener are known in the art. Common synthetic thickeners are polyisobutylene (PIB), VI improvers, such as poly(methyl)methacrylate, olefin copolymers and the like, and polyalphaolefins (PAO) having a high kinematic viscosity. An example of a PAO thickener is PAO 100, i.e. a PAO having a Vk,100 of about 100 cSt. Such high viscosity PAO is used to obtain the multigrade properties and the desired viscosity, whilst maintaining thermal and oxidation stability. In addition to such PAO a low viscosity ester is normally used to improve the solubility and compatibility of the additives used, to enhance thermal stability and oxidation stability and to impart the desired low temperature viscosity to the gear oil formulation. An EP/AW additive package is applied to avoid wear of the gear wheels. Finally, a low viscosity (i.e. Vk,100 of 4-10 cSt) PAO, also denoted as PAO 4 to PAO 10, and/or a mineral oil having a high viscosity index (VI) is normally present as a base fluid. In case a fully synthetic multigrade gear oil is desired, a low viscosity PAO is used.

It has been found, however, that although the current synthetic multigrade gear oils containing a synthetic thickener perform satisfactorily in a number of demanding applications, there is still a need for improvement to cope with the increasing requirements of modern gear oils such as for heavy duty commercial vehicles and for pessenger cars with long drain intervals or filled for life systems. It is an object of the present invention to provide a multigrade gear oil formulation having an improved performance, particularly in gear boxes for heavy duty vehicles, and which also can be fully synthetic, although the latter is not specifically required.

It has been found that by using the complex esters as described hereinbefore as a thickener the above objects can be realised.

Accordingly, the present invention also relates to a multigrade gear oil formulation comprising:

PCT/EP98/06145 WO 99/16849

5-45 pbw of the complex ester as described hereinbefore as a thickener, (a)

- 5-45 pbw of an ester having a kinematic viscosity at 100 C of 2-10 cSt, (b)
- 5-60 pbw of a mineral oil having a VI of at least 90 and/or a (c) polyalphaolefin having a kinematic viscosity at 100 C of 4-10 cSt, and
- 5 (d) 5-15 pbw of the usual gear oil additives, the sum of the amounts of the components (a) to (d) being 100 pbw.

10

30

Components (b), (c) and (d) can be any ester, mineral oil and/or polyalphaolefin and additives known to be useful or already used in multigrade gear oil formulations.

Component (b), the low viscosity ester, may be any ester suitable for improving additive solubility and compatibility as well as for improving thermal and oxiation stability and for imparting the desired low temperature viscosity to the gear oil formulation. Preferably, component (b) is an ester of a neopentyl polyol, suitably trimethylolpropane, with at least one aliphatic, saturated monocarboxylic acid having 6 15 to 12 carbon atoms. An example of such ester is commercially available under the trade name PRIOLUBE 3970.

Component (c) may be a mineral oil or a PAO, which should have a VI of at least 90. It is, however, preferred to use a PAO, particularly PAO 6 and PAO 8.

Component (d) may be any available gear oil EP/AW additive package 20 known to be useful in automotive and industrial gear oil formulations.

The complex esters may be produced in a batch or continuous process. The invention further provides a process for the manufacture of a complex ester which comprises reacting at least one polyfunctional alcohol, at least one polyfunctional carboxylic acid and a chain stopping agent, wherein

- 25 (a) the polyfunctional alcohol is a hindered or non-hindered, aliphatic polvol,
 - the polyfunctional carboxylic acid comprises an aliphatic dicarboxylic (b) acid containing from 9 to 18 carbon atoms, dimerised and/or trimerised fatty acids or mixtures thereof, with the proviso that dimerised and trimerised fatty acids do not constitute more than 80% by weight, preferably not more than 50% by weight, of the total amount of polyfunctional carboxylic acid used,

the chain stopping agent comprises either an aliphatic monocarboxylic acid selected from the group consisting of straight chain saturated acids containing from 7 to 22, preferably from 7 to 14, carbon atoms, branched saturated acids containing from 7 to 24 carbon atoms, straight or branched unsaturated acids containing from 16 to 24 carbon atoms and mixtures thereof or at least one aliphatic, straight or branched, saturated or unsaturated, monofunctional alcohol containing at least 14 carbon atoms, and preferably not having more than 24 carbon atoms, and

(d) the complex ester has a kinematic viscosity at 100 C (Vk,₁₀₀) of from 30 to 1000 cSt, preferably from 30 to 200 cSt.

The invention is further illustrated by the following examples without limiting the scope of the invention to these examples.

Example 1

5

10

Two complex esters were prepared by esterification of the following mixtures:

	Ester A:	Ester B:
	19 pbw trimethylolpropane	18 pbw trimethylolpropane
	22 pbw dodecanedioic acid	18 pbw decanedioic acid
20	59 pbw isostearic acid	6 pbw dimer acid
	~	58 pbw isostearic acid

Ester A had a Vk_{100} of 117,0 cSt and a Vk_{40} of 1360 cSt. Ester B had a Vk_{100} of 121,6 cSt and a Vk_{40} of 1445 cSt.

25

Each complex esters was formulated into a gear oil formulation having the following composition:

	30.0 pbw	complex ester A or B
	35.8 pbw	PAO 8
30	25.0 pbw	PRIOLUBE 3970
	9.2 pbw	HITEC 381 (trade mark), a sulphur-phosporus-
		containing EP/AW additive package sold by Ethyl Corp.

The formulation containing complex ester A is denoted as Formulation A, the formulation containing complex ester B as Formulation B.

Both Formulations A and B were subjected to a severe screening test being the CEC L-48-A-95 (A) oxidation test, also known as the GFC test. This test is widely known and used in the industry to measure the oxidation stability of lubricating oils used in automotive transmissions by artificial ageing.

In the test samples are subjected to oxidation conditions by heating to a temperature of 160 C and by passing air through the samples at a flow rate of 10 litres per hour during a period of 192 hours. However, to increase test severity and to demonstrate the excellent properties of complex esters A and B, the test duration was extended to 300 hours.

The results are indicated in Table 1.

Comparative Example 1

5

10

25

30

A gear oil formulation (Formulation C) similar to Formulations A and B, only comprising 30.0 pbw of PAO 100 as a thickener instead of a complex ester, was also subjected to the severe screening test of Example 1.

The results are indicated in Table 1

20 Table 1 Gear oil formulation performance

ForeSulation	А	В	С
Change in Vk,,,,, (%)	9	15	32
Change in Vk, ₄₀ (%)	16	24	84
Pentane insolubles (%)	0.11	0.15	0.65
Toluene insolubles (%)	0.11	0.12	0.59

From Table 1 it can be seen that formulations A and B show a significantly better performance than Formulation C. both with regard to change of viscosity and insolubles, which indicate that the oxidation stability of Formulations A and B is better than that of Formulation C. During oxidation, namely, viscosity changes and insolubles are formed. The smaller the change in viscosity and the less insolubles are formed, the better the oxidation stability.

Example 2

25

Two other complex esters were prepared by esterification of the following mixtures:

Ester D: Ester E:

5 13 pbw pentaerythritol 13 pbw pentaerythritol

9 pbw decanedioic acid 14 pbw dodecanedioic acid

78 pbw isostearic acid 73 pbw isostearic acid

Ester D had a Vk_{100} of 54,0 cSt and a Vk_{40} of 471 cSt.

10 Ester E had a Vk_{100} of 93,5 cSt and a Vk_{40} of 1105 cSt.

Ester D and Ester E were subjected to biodegradation tests according to OECD-Guideline 301 B (modified Sturm test). The test is based on the measurement of CO₂ evolution and is a well-known and widely accepted test to measure ultimate biodegradability. Ultimate biodegradability relates to the conversion of the parent molecule to simple molecules such as carbon dioxide, water, inorganic salts and new micro-organisms.

After the prescribed test period of 28 days. Ester D was biodegraded to an extend of 65% and Ester E to an extend of 63%. Based on the OECD 301 B ready

biodegradability threshold of >/= 60% after 28 days, both Ester D and Ester E may be termed readily biodegradable.

Their ready biodegradability make such esters as Ester D and Ester E well suitable for application in biodegradable greases, biodegradable chain oils, biodegradable hydraulic fluids, biodegradable industrial gear oils and the like. For these applications, the esters may be used as such and/or in combination with other readily biodegradable base fluids such as other complex esters, non-complex esters, polyalphaolefins of both suitable viscosity and biodegradability and certain mineral oil type of base fluids. The formulations containing these product may also contain suitable additives such as antioxidants, anti-wear/extreme pressure additives, metal deactivators, anticorrosion

additives, antifoamants, friction modifiers and the like as known in the art.

Example 3

Two other complex esters were prepared by esterification of the following mixtures:

Ester F: Ester G:

5 32 pbw neopentylglycol 35 pbw dipropylene glycol

48 pbw decanedioic acid 38 pbw dodecanedioic acid

11 pbw octanoic acid 15 pbw octanoic acid

9 pbw decanoic acid 12 pbw decanoic acid

Ester F had a Vk, of 45,4 cSt and a Vk, of 402 cSt.
 Ester G had a Vk, of 31,8 cSt and a Vk, of 231 cSt.

Ester F and Ester G have a particular polar character as a result of the presence of a high amount of ester groups which results in excellent lubricity, in particular in relation to non-polar base fluids such as mineral oil and/or synthetic hydrocarbons and/or less polar esters. Therefor, such esters as Ester F and Ester G are suitable for use as a base fluid component and/or additive in engine oils to reduce the internal friction of those engines. For this application, the esters may be used as such and/or in combination with other base fluids such as non-complex esters, polyalphaolefins and mineral oil type of base fluids. The formulations containing these product may also contain suitable additives such as detergents, dispersants, antioxidants, anti-wear/extreme pressure additives, metal deactivators, anticorrosion additives, antifoamants, friction modifiers and the like as known in the art.

25 Example 4

15

20

An other complex ester was prepared by esterification of the following mixtures:

Ester H:

26 pbw pentaerythritol

30 23 pbw hexanedioic acid

51 pbw hexanoic acid

Ester H had a Vk,100 of 217 cSt and a Vk,40 of 3265 cSt.

Ester H has a very high affinity to metal surfaces due to the presence of a very high amount of ester groups. Therefor, such an ester is suitable for use as a base fluid component and/or additive in metal working oils to improve the lubricity of the formulation, thereby improving the metal working process. The esters may be used in combination with other base fluids such as other esters, polyalphaolefins and mineral oil type of base fluids. The formulations containing these product may also contain suitable additives such as antioxidants, anti-wear/extreme pressure additives, metal deactivators, anticorrosion additives, antifoamants and the like as known in the art.

Example 5

An other complex ester was prepared by esterification of the following mixtures:

15 Ester I:

5

10

23 pbw dipentaerythritol

8 pbw hexanedioic acid

38 pbw octanoic acid

31 pbw decanoic acid

20

Ester I had a Vk,100 of 35,5 cSt and a Vk,40 of 329 cSt.

Owing to its high oxidation stability and good lubricity due to the presence of a polar ester groups such an ester is suitable for use as a base fluid component and/or additive for compressor oils and for metal rolling oils. The esters may be used in combination with other base fluids such as other esters, polyalphaolefins and mineral oil type of base fluids. The formulations containing these product may also contain suitable additives such as antioxidants, anti-wear/extreme pressure additives, metal deactivators, anticorrosion additives, antifoamants and the like as known in the art.

30

25

Claims

10

15

20

1. A complex ester obtainable by an esterification reaction between at least one polyfunctional alcohol and at least one polyfunctional carboxylic acid and a chain stopping agent, wherein

- 5 (a) the polyfunctional alcohol is a hindered or non-hindered, aliphatic polyol,
 - (b) the polyfunctional carboxylic acid comprises an aliphatic dicarboxylic acid containing from 9 to 18 carbon atoms, dimerised and/or trimerised fatty acids or mixtures thereof, with the proviso that dimerised and trimerised fatty acids do not constitute more than 80% by weight of the total amount of polyfunctional carboxylic acid used,
 - the chain stopping agent comprises either an aliphatic monocarboxylic acid selected from the group consisting of straight chain saturated acids containing from 7 to 22 carbon atoms, branched saturated acids containing from 7 to 24 carbon atoms, straight or branched unsaturated acids containing from 16 to 24 carbon atoms and mixtures thereof or at least one aliphatic, straight or branched, saturated or unsaturated, monofunctional alcohol containing at least 14 carbon atoms, and
 - (d) the complex ester has a kinematic viscosity at 100 C (Vk,₁₀₆) of from 30 to 1000 cSt.
 - 2. Complex ester according to claim 1, wherein the polyfunctional alcohol is a hindered polyol, preferably a neopentyl polyol.
 - 3. Complex ester according to claim 2, wherein the neopentyl polyol is trimethylolpropane or pentaerythritol.
- 4. Complex ester according to any one of the preceding claims, wherein the aliphatic dicarboxylic acid has from 9 to 12 carbon atoms.
 - 5. Complex ester according to any one of the preceding claims, wherein the chain stopping agent is isostearic acid.
- 6. Complex ester according to any one of the preceding claims, wherein the complex esters have a kinematic viscosity at 100 C of from 100 to 140 cSt.

7. Complex ester according to any one of the preceding claims, wherein the polyol, polyfunctional carboxylic acid and chain stopping agent are used in the following amounts:

15-20 pbw of polyol,

- 5 20-25 pbw polyfunctional carboxylic acid and 55-65 pbw chain stopping agent.
 - 8. A functional fluid composition comprising a complex ester as defined in any one of claims 1 to 7.
- 9. A functional fluid composition according to claim 8 which further comprises an additive package containing a sulphur and/or phosphorus-containing extreme pressure and/or anti-wear compound in a weight ratio of complex ester to additive package of from 1:3 to 9:1.
 - 10. Use of the complex ester according to any one of claims 1 to 7 as a functional fluid.
- 15 11. Use of the complex ester according to any one of claims 1 to 7 as an additive and/or as a base fluid and/or as a thickener in a functional fluid composition.
 - Use according to claim 10 or 11 in which the functional fluid or functional fluid composition is a lubricating oil, a transmission oil, a gear oil, an axle oil, an automatic transmission fluid, a hydraulic fluid, a four-stroke oil, a fuel additive, a compressor oil a grease a chain oil or a lubrication oil for a transmission.
- compressor oil, a grease, a chain oil or a lubricating oil for metal working or metal reling applications.
 - 13. A process for the manufacture of a complex ester which comprises reacting at least one polyfunctional alcohol, at least one polyfunctional carboxylic acid and a chain stopping agent, wherein
- 25 (a) the polyfunctional alcohol is a hindered or non-hindered, aliphatic polyol,

30

(b) the polyfunctional carboxylic acid comprises an aliphatic dicarboxylic acid containing from 9 to 18 carbon atoms, dimerised and/or trimerised fatty acids or mixtures thereof, with the proviso that dimerised and trimerised fatty acids do not constitute more than 80% by weight, preferably not more than 50% by weight, of the total amount of polyfunctional carboxylic acid used,

(c) the chain stopping agent comprises either an aliphatic monocarboxylic acid selected from the group consisting of straight chain saturated acids containing from 7 to 22, preferably from 7 to 14, carbon atoms, branched saturated acids containing from 7 to 24 carbon atoms, straight or branched unsaturated acids containing from 16 to 24 carbon atoms and mixtures thereof or at least one aliphatic, straight or branched, saturated or unsaturated, monofunctional alcohol containing at least 14 carbon atoms, and preferably not having more than 24 carbon atoms, and

- (d) the complex ester has a kinematic viscosity at 100 C (Vk,₁₀₀) of from 30 to 1000 cSt, preferably from 30 to 200 cSt.
 - 14. Multigrade gear oil formulation comprising:
- (a) 5-45 pbw of at least one complex ester according to any one of claims 1-7 as a thickener,
- (b) 5-45 pbw of an ester having a kinematic viscosity at 100 C of 2-10 cSt,
- 15 (c) 5-60 pbw of a mineral oil having a VI of at least 90 and/or a polyalphaolefin having a kinematic viscosity at 100 C of 4-10 cSt, and
 - (d) 5-15 pbw of the usual gear oil additives, the sum of the amounts of the components (a) to (d) being 100 pbw.
- 15. Gear oil formulation according to claim 14, wherein the low viscosity ester is 20 an ester of a neopentyl polyol, preferably trimethylolpropane, with at least one aliphatic, saturated monocarboxylic acid having 6 to 12 carbon atoms.
 - 16. Gear oil formulation according to claim 14 or 15, wherein component (c) is a polyalphaolefin selected from PAO 6 and PAO 8.

25

5

10

30

INTERNATIONAL SEARCH REPORT

International Application No PCT/EP 98/06145

PCT/EP 98/06145 CLASSIFICATION OF SUBJECT MATTER C10M105/42 C10M129/78 C10M105/44 C10M105/46 C10M129/80 C10M129/82 C10M169/04 C10M171/00 C10L1/18 C07C69/34 C07C69/50 C07C69/593 C07C69/604 According to International Patent Classification (IPC) or to both national classification and IPC Minimum documentation searched (classification system followed by classification symbols) IPC 6 C10M C10L Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Electronic data base consulted during the international search (name of data base and, where practical, search terms used) C. DOCUMENTS CONSIDERED TO BE RELEVANT Category: Citation of document, with indication, where appropriate, of the relevant passages Relevant to claim No X EP 0 415 778 A (KAO CORP) 6 March 1991 1-4.6-13 see page 3. line 3 - page 4. line 4 see page 5. line 15 - line 49 χ GB 1 460 665 A (CIBA GEIGY AG) 1-13 6 January 1977 see page 1. line 20 - page 2, line 65 X WO 94 01516 A (HENKEL KGAA :BONGARDT FRANK 1-13 (DE); WINDGES NICOLE (DE)) 20 January 1994 see page 2, paragraph 5 - page 5. paragraph 4 see page 7. paragraph 2 - page 8. paragraph 1 χ EP 0 535 990 A (NIPPON OIL CO LTD) 1 - 137 April 1993 see page 2. line 22 - page 3. line 58 Further documents are listed in the continuation of box C X Patent family memoers are listed in annex. Special categories of cited documents T" later document published after the international filling date or priority date and not in conflict with the application out cited to understand the principle or theory underlying the "A" document defining the general state of the lart which is not considered to be of particular relevance "E" earlier document but published on or after the international "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to nling date "L" accument which may threw doubts on phority iclaim(s) or involve an inventive step when the document is taken alone which is cited to establish the publication date of another document of particular relevance; the claimed invention offation or other special reason (as specified) cannot be considered to involve an inventive, step when the "O" secument referring to an oral disclosure, use, exhibition or document is combined with one or more other, such docu ments, such combination being obvious to a person skilled document published prior to the international filing date but later than the priority date claimed. in the art. "3" document member of the same patent family Date of the actual completion of the international search Date of mailing of the international search report 17 February 1999 02/03/1999 Name and mailing address of the ISA Authorized officer European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl. Rotsaert. L Fax: (+31-70) 340-3016

INTERNATIONAL SEARCH REPORT

Inte ional Application No
PCT/EP 98/06145

		PC1/EP 98/06145
C.(Continu	ation) DOCUMENTS CONSIDERED TO BE RELEVANT	
Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No
X	PATENT ABSTRACTS OF JAPAN vol. 014, no. 208 (C-0714). 27 April 1990 & JP 02 045595 A (KAO CORP), 15 February 1990 see abstract	1-13
X	PATENT ABSTRACTS OF JAPAN vol. 095, no. 005, 30 June 1995 & JP 07 034080 A (KAO CORP), 3 February 1995 see abstract	1-13
X	DATABASE WPI Section Ch. Week 7850 Derwent Publications Ltd London, GB: Class A17. AN 78-90446A XP002093831 & JP 53 127970 A (NIPPON OILS & FATS CO LTD), 8 November 1978 see abstract	1-16
Α	EP 0 578 435 A (ETHYL PETROLEUM ADDITIVES LTD) 12 January 1994 see page 3, line 54 - page 4. line 16	14-16
A	DE 26 20 645 A (INST FRANCAIS DU PETROL;RHONE POULENC IND (FR)) 2 December 1976 cited in the application see the whole document	1,13
	-	

1

INTERNATIONAL SEARCH REPORT

Information on patent family members

Int tional Application No PCT/EP 98/06145

			70,002.0				
Patent document cited in search repo	rt	Publication date		Patent family member(s)	Publication date		
EP 0415778	A	06-03-1991	JP JP DE US	2801703 B 3179091 A 69002209 T 5096606 A	21-09-1998 05-08-1991 28-10-1991 17-03-1992		
GB 1460665	Α	06-01-1977	NON	E			
WO 9401516	А	20-01-1994	DE AT DE EP ES JP US	4222341 A 138681 T 59302760 D 0649457 A 2087751 T 7508783 T 5503762 A	13-01-1994 15-06-1996 04-07-1996 26-04-1995 16-07-1996 28-09-1995 02-04-1996		
EP 0535990	Α	07-04-1993	JP AT CA DE	5098276 A 147096 T 2079512 A 69216346 D	20-04-1993 15-01-1997 05-04-1993 13-02-1997		
EP 0578435	Α	12-01-1994	CA DE DE JP	2099314 A 69314554 D 69314554 T 6179887 A	10-01-1994 20-11-1997 19-02-1998 28-06-1994		
DE 2620645	A	02-12-1976	FR BE CH	2311088 A 841640 A 613471 A	10-12-1976 10-11-1976 28-09-1979		

			4	٠	*
					:
					,
-					
					••